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(54) **COMPACT AUTOMATED CELL COUNTER**

KOMPAKTER AUTOMATISierter ZELLZÄHLER

COMPTEUR DE CELLULES AUTOMATISÉ COMPACT

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Description

BACKGROUND OF THE INVENTION

Field of the Invention

[0001] This invention lies in the field of hemocytometry and systems in general for the counting of biological cells suspended in fluids. The focus of this invention is on automated cell counting systems.

Description of the Prior Art

[0002] Cell counting is of interest in a variety of clinical and research procedures, including the counting of leukocytes and erythrocytes, which is of value in the diagnosis of various diseases or abnormal conditions and in the monitoring of patients that are undergoing treatment for such diseases or conditions. Cells can be counted manually by placing a known dilution of a sample between optically clear plates that are sufficiently close to each other (typically with a spacing on the order of 100 μm) to form the cells into a single layer, magnifying an area of the layer of designated dimensions to a known magnification, and counting the cells in the magnified area through a microscope. Manual cell counters often include a grid inscribed in the counting area to lessen the burden on the user. A description of such a grid and the procedure for its use is found in Qiu, J., United States Patent No. US 7,329,537 B2, issued February 12, 2008, "Micro-Pattern Embedded Plastic Optical Film Device for Cell-Based Assays." Regardless of how it is done, manual cell counting is tedious and highly vulnerable to user error. Counting is commonly aided by using a high dilution of the sample to lessen the number of cells in the counting area, but the accuracy of the counting declines with every decrease in the proportion of cells that are counted.

[0003] Automation of cell counting procedures has been made possible by the use of digital imaging systems. An example of such a system is ImageJ, a Java-based image processing program developed at the National Institutes of Health and reported by Collins, T.J., "ImageJ for microscopy," *BioTechniques* 43 (1 Suppl.): 25-30 (July 2007). The use of ImageJ in hematology systems is reported by Gering, T.E., and C. Atkinson, "A rapid method for counting nucleated erythrocytes on stained blood smears by digital image analysis," *1 Parasitol.* 90(4): 879-81 (2004). Further disclosures of automated cell counting are Chang, J.K., *et al.*, United States Patent No. US 7,411,680 B2, issued August 12, 2008, "Device for Counting Micro Particles," and Chang, J.K., *et al.*, United States Patent Application Publication No. US 2006/0223165 A1, published October 5, 2006, "Device for Counting Cells and Method for Manufacturing the Same." Other examples of prior art devices are shown in : US2004/170312, US2008/088952, US2003/026762, US2004/246479, US5754291, US2005/023439 or WO98/50577.

[0004] Automated cell counting systems themselves contain an inherent statistical uncertainty due to what is commonly referred to as "sampling error," which refers to the error inherent in selecting the area in which the automated counting is performed. One of the limitations of automated cell counters that are currently available is that due to the limitations of the optical components in the instruments, the area in which cells are counted is of limited size compared to the entire area occupied by the sample. Since this limits the number of cells accordingly, and the error increases with every decrease in the number of cells being counted, the typical instrument of the prior art is constructed with a long optical path or a large footprint (the surface area on a laboratory bench that the instrument consumes), or both, to achieve an acceptable level of accuracy. This presents disadvantages to the user, particularly when the instrument is to be used in a cell culture hood.

SUMMARY OF THE INVENTION

[0005] Disclosed herein is a fully self-contained instrument for highly accurate cell counting with minimal user intervention as well as a relatively small footprint and limited height. A cell suspension is placed in a consumable sample vessel whose size and dimensions can vary widely, one convenient example of which is a vessel whose outer dimensions are similar to those of a microscope slide. The vessel can thus be similar in construction and dimensions to the vessel described in US 2006/0223165 A1 referenced above, with at least one flat, shallow internal chamber bounded on the top and bottom by flat, optically clear windows, which can be plastic sheets, whose spacing is close enough that most of the cells of the sample form a layer that is one cell deep. Appropriate inlet and vent ports can be included in the vessel to allow the chamber to be easily and completely filled with the sample. The vessel is then placed in the instrument where it intersects a linear optical path.

In particular the invention provides an automated cell counter comprising optical components, a sample mount for receiving a sample slide, and a housing retaining said optical components and sample mount, said optical components comprising a light source, a digital imaging sensor, and lenses positioned to direct light from said light source to said digital imaging sensor through said sample mount, said lenses comprising a negative lens adjacent to said digital imaging sensor, and two achromat lenses of an achromat lens pair separated by an aperture, the ratio of the focal lengths of said two achromat lenses being within a range of 1.5 to 6, said optical components arranged in a linear optical path and arranged such that when a sample slide containing a cell suspension is mounted in said sample mount, an image of said cell suspension is projected onto said digital imaging sensor, said digital imaging sensor comprising at least about 4,000,000 pixels and imaging a field of view at said sample mount of at least 3 mm^2 , each pixel being $2 \times 2 \mu\text{m}$

or less in area, wherein said optical path is vertically oriented and said housing is mounted to a support base that is at least as wide as said housing, said housing and support base each occupying lateral areas of not more than 300 cm² and said optical path having a total height of not more than 20 cm.

The term "linear" as used herein denotes a path with no turns or other changes in direction of the light beams other than those caused by lenses. The vessel enters the instrument through a slot at a designated height in the optical path, and as described below in greater detail, the instrument in certain embodiments of the invention contains features that automatically adjust the height of the sample for purposes of focusing the sample image. Certain embodiments contain features that cause all instrument functions to begin operation upon the insertion of the sample vessel into the instrument.

The invention further provides a method for counting cells in a cell suspension, said method comprising: (a) placing an aliquot of said cell suspension in an internal chamber of a sample slide to form a film of said suspension in said chamber, (b) creating a digital image of a field of view of at least 3 mm² within said film on the automated cell counter as defined above and (c) digitally counting said cells in said digital image.

BRIEF DESCRIPTION OF THE DRAWINGS

[0006]

FIG. 1 is a perspective view of a cell counting instrument representing an example of an implementation of the concepts disclosed herein.

FIG. 2 is a diagram of the optical components of the instrument of FIG. 1.

FIG. 3 is a perspective view of optical components in the interior of the instrument of FIG. 1.

FIG. 4 is an exploded view, in perspective, of two plates constituting a sample slide for use in the instrument of FIG. 1.

FIG. 5A is a view of the upper surface of the upper plate of the sample slide of FIG. 4. fig. 5B is a view of the lower surface of the upper plate of the sample slide of FIG. 4.

DETAILED DESCRIPTION

[0007] The upper and lower optical windows between which the cell suspension is retained inside the sample vessel are close enough that the retained suspension is a thin film whose lateral dimensions, *i.e.*, its exposed length and width, are at least an order of magnitude greater than its thickness. The entire exposed area (*i.e.*, lateral dimensions) of the sample chamber or a laterally dimen-

sioned portion thereof serves as a field of view that is projected onto a digital imaging sensor that contains at least 4,000,000 (four million) pixels, or in certain embodiments from 4,000,000 to 10,000,000 pixels, with each pixel being no greater than 2 x 2 μm (4 μm²) in size, or from 0.5 x 0.5 μm (0.25 μm²) to 2 x 2 μm (4 μm²) in certain embodiments, and in certain of the latter from 1 x 1 μm (1 μm²) to 2 x 2 μm (4 μm²). The field of view imaged by the sensor is at least 3 mm², and often from 3 mm² to 10 mm². A complementary metal oxide semiconductor (CMOS) is one example of a digital imaging sensor useful for this purpose. Examples of CMOS sensors meeting these parameters are the OV5620 and OV5632 color imagers available from OmniVision, Santa Clara, California, USA. Other examples are available from Aptina Imaging, a division of Micron Technology, Inc., of San Jose, California, USA. A color digital imaging sensor can also be used. Image processing to count the cells in the image generated by the CMOS sensor can be achieved by known digital counting methods, such as those mentioned above.

[0008] The image of the sample chamber can be magnified along the optical path by a magnification that is often within a range of from 1.5 to 6, or a range of from 1.5 to 3, with a magnification of about 2 as an example. This can be achieved by a two-lens achromat assembly. An example of such a lens assembly is a lens of 35-mm focal length closest to the sample, a lens of 60-mm focal length closest to the sensor, and an aperture between the two lenses. The distance between the lens nearest the sample and the sample itself in this example is thus 35 mm, and the distance between the lens nearest the sensor and the imager itself is 60 mm. The magnification of the system is the ratio of the focal lengths of the two lenses, which in this case is 60 mm/35 mm = 1.7. The two lenses can each for example be 12.5 mm in diameter, and the aperture can be 6mm in diameter. Lenses of other diameters and focal lengths that will produce the same or approximately the same results will be readily apparent to those skilled in the art. The footprint of the instrument is defined as the area projected by the larger of the instrument and its support base on a plane perpendicular to the optical path. As noted above, the instrument can be constructed with a small footprint, particularly one that is less than 300 cm² in area.

[0009] When a flat digital imaging sensor is used, a negative lens can be positioned below the sensor to intercept the optical signal immediately and to correct the focus field curvature of the achromat lens pair. This type of field curvature is common in optical systems and is also referred to as Petzval curvature. In an illustrative embodiment, a 6 mm-diameter lens with a minus-18 mm focal length is used. The lens thickness can vary but is optimally selected to correct the curvature without substantially reducing the field of view.

[0010] Illumination of the sample can be achieved with a conventional light source at the base of the instrument and a collimating lens between the light source and the

sample. With these components the sample is illuminated by trans-illumination without a diffuser. A preferred light source is a single white light-emitting diode (LED) with a fluorescent coating. An example of such a component is LUXEON® Rebel White, part no. LXML-PWN1-0050, available from Philips Lumileds Lighting Company, San Jose, California, USA. An example of a collimating lens is one that is 9 mm in diameter with a focal length of 18 mm. With these dimensions and those of the preceding paragraphs, an instrument can be constructed with the achromat lens pair approximately 35 mm above the sample, and the sensor approximately 60 mm above the achromat lens pair. With an achromat lens pair having a thickness of approximately 13 mm, the total distance between the sample and the sensor can be as little as 108 mm. In general, the optical path of the instrument, *i.e.*, defined herein as the arrangement of the components extending from the light source to the CMOS or other digital imaging sensor, can be 20 cm or less in height. In preferred instruments within the scope of this invention, the optical components are mounted to the housing interior in a floating manner using compliant counts, to avoid damage to, or misalignment of, the optical system upon jolts to the instrument, such as might occur when the instrument is dropped or mishandled, or collides with another instrument or piece of equipment.

[0011] As noted above, the sample vessel, which will be referred to henceforth as a sample slide in view of its similarity in size and shape to a microscope slide, is received in the instrument through a slot that is positioned at a location along the optical path that is at a distance from the nearest lens of the achromat lens pair equal to the focal length of the lens. In its preferred embodiments, the instrument as a whole is 30 cm or less in height, and the use of a digital imaging sensor as described above that employs a large number of pixels of the small sizes indicated permits the instrument to be constructed with the slot at a sufficient height to allow the user to comfortably insert the slot by hand, *i.e.*, clearing the user's hand from the table on which the instrument rests. The slot can thus be 60 mm or more from the base of the instrument, and preferably 70-80 mm from the base.

[0012] In preferred embodiments of the invention, the instrument provides autofocusing of the sample image by automatically adjusting the height of the slide following its insertion. One means of autofocusing involves the use of an image processor chip that provides an output of image contrast within an array of zones across the image from the sensor. An example of such a chip is the Freescale Semiconductor MC9328MX21, available from Keil™ - an ARM Company, Plano, Texas, USA; other examples will be apparent to those skilled in the art. The sum of the absolute differences of adjacent green pixels in a particular zone of the sensor array can be used as the image contrast value, and optimum focus is achieved when the image contrast value is at a maximum. The focus can then be adjusted by a geared motor connected to the slide mount within the receiving slot, *i.e.*, the motor

when rotated will move the slide mount up or down to change the focus of the image. The contrast value is detected at various positions of the motor which is then returned to the position producing the highest contrast value. In many embodiments of the instrument, this autofocusing can occur in 15 s or less.

[0013] An accessory that can be supplied with the instrument is a standard slide for quality control, such as verifying the accuracy of counting live and dead cells and the ability of the instrument to focus properly. The standard slide can have the same external dimensions as a sample slide, but instead of a sample chamber(s), the standard can have an array of dark-colored spots and rings printed on it, the spots simulating dead cells and detected as such in the digital imaging sensor and the rings simulating live cells and detected as such in the digital imaging sensor.

[0014] In certain embodiments of the concepts described herein, the functions performed by the instrument, including autofocusing and cell counting, are initiated by the simple insertion of the sample slide. This initiation can be achieved by the inclusion of a non-contact optical reflection sensor located within the slot or on the slide mount within the slot.

An example of a suitable sensor is one that emits an infra-red beam and detects objects within approximately one millimeter of the sensor aperture by detecting a reflected signal from the beam. The reflected signal will rise to a maximum level when the slide is fully inserted, and the high signal will initiate the autofocusing and cell counting mechanisms. One example of a sensor that can serve this purpose is the QRE1113 Reflective Object Sensor, available from Fairchild Semiconductor Corporation, San Jose, California, USA. Other examples will be apparent to those skilled in the art.

[0015] A further feature that can be included in instruments embodying the features described herein is the automatic detection of cells in the sample that are stained with a vital stain. A vital stain is one that preferentially stains dead cells, and the differentiation between cells stained with such a stain and those that are not is achieved by the use of differently colored pixels. Trypan blue is one example of a vital stain; eosin and propidium iodide are other examples. Trypan blue transmits blue light and attenuates red light, and by comparing the intensities of blue and red pixels in the image sensor, the instrument can determine whether cells stained with a vital stain are present. Other dyes will afford similar color distinctions as appropriate to the dyes themselves. Image processing chips that incorporate this automatic detection feature include those referenced above and are readily available. The instrument can be programmed to eliminate any possible undercounting of viable cells and thereby detect viable cells to a particularly high degree of accuracy by focusing on two or more planes. The contrast between live cells and dead cells can be increased further by using optical filters to control the illumination bandwidth, or by selecting a spectrally narrow light

source, such as an LED of a particular color instead of white. For example, a 585 nm optical filter with about 20 nm bandwidth can be used to match the illumination to the peak absorption wavelength of the Trypan blue dye, whose peak absorption is 586 nm. The dead cells will appear darker when the sample is illuminated through this filter.

[0016] In preferred instruments within the scope of this invention, all functions that contribute to the obtainment of a cell count in the sample are contained within the instrument housing, and the full Operation of the instrument can thus be achieved without the use of an external machine or computer. Included among these functions are the automatic focusing by varying the height of the sample slide to find the best focal plane to discriminate cells from background, the determination of whether the sample has been stained with Trypan blue or other vital stain, a multi-focal plane analysis when a vital stain is detected so that each cell is scored on multiple focal planes to prevent undercounting of live cells, an integrated dilution counter to determine the volume of a cell suspension to use, the ability to produce a visual image of the cells on the display at the option of the user and to zoom in for a detailed visual inspection of the cells, and the ability and user option to export the results to a USB flash drive or to a thermal printer or other external printer. All of these functions can be initiated by the simple insertion of the sample slide by way of the non-contact optical reflection sensor described above, and in many cases, the execution of these functions is completed in 30 s or less.

[0017] The Figures hereto depict an instrument that contains many of the features described above and serves as one example of an implementation of the concepts described herein.

[0018] FIG. 1 depicts an automated cell counter instrument 11 in its upright position as it would be used on a laboratory bench. The visible parts of the instrument are a housing 12, a support base 13, a display screen 14, a control panel 15, and a slot 16 for insertion of a sample slide 17. The display screen shows the progress of the cell counting analysis, identifies the functions of the instrument as they are being performed, and offers options to the user for various functions and for showing an image of the cells in the sample slide.

[0019] FIG. 2 depicts components of the optical path in the interior of the instrument of FIG. 1 with the sample slide 17 having been positioned in the optical path. The sample slide 17 is horizontal and resides above an LED board 22 serving as the light source. A collimating lens 23 renders the light rays from the LED parallel as they approach the sample slide. The achromat lens pair 24 is positioned between the sample slide 17 and the sensor 25. The two lenses 26, 27 of the achromat lens pair are separated by an aperture 28. A field flattening lens 29 is positioned immediately below the sensor 25.

[0020] FIG. 3 depicts the main optics assembly, showing the slide mount 31 with the sample slide 17 partially

inserted, the LED board 22, the illumination (collimating) lens 23, the geared motor 32 that adjusts the slide height to focus the image, and an imaging lens tube 33 terminating in a fitting 34 to receive the CMOS sensor board. Also shown in the Figure is the main printed circuit board 35 that controls the functions of the instrument and includes a motor drive chip to control the motor 32. The board 35 resides within the housing and the position of the board in the Figure reflects its position relative to the optics assembly.

[0021] A sample slide for use in the instrument of the preceding Figures is shown in FIGS. 4, 5A, and 5B. The view in FIG. 4 is a perspective view, and the slide 17 is formed of two plates 42, 43 bonded together but shown separated in the Figure. The slide contains two sample chambers, as indicated by the indicia "A" and "B," respectively, separated from each other lengthwise along the slide and laterally offset from each other. The areas 44, 45 of the lower plate 43 that form the bottom surfaces of the sample chambers are made of optically transparent material, as are the corresponding areas of the upper plate 42 that are directly above these areas on the lower plate and form the upper surfaces of sample chambers. The lower plate 43 in this embodiment is thicker than the upper plate 42 to provide rigidity to the slide, and the relative thinness of the upper plate 42 permits the upper window of each sample chamber to be thinner than the lower window, and indeed as thin as possible to achieve a highly focused image in the CMOS sensor. Each sample chamber is thus offset from the center plane of the slide and closer to the upper plate 42 than to the lower plate 43.

[0022] FIGS. 5A and 5B are planar views of the top surface 51 and bottom surface 52, respectively, of the upper plate 42, the bottom surface 52 being the surface that is bonded to the lower plate 43. Each sample chamber is defined by a recess 53 (FIG. 5B) in the bottom surface of the upper plate, which further reduces the thickness of the area forming the optically clear window at the top of each sample chamber. In one example of the dimensions of the slide, the thickness of the upper plate in areas other than the recess 53 is 0.65 mm and the thickness of the lower plate is 1.00 mm, for a total slide thickness of 1.65 mm. The recess 53 is 0.100 mm in depth, which thus forms a sample chamber that is 0.100 mm in depth, a standard sample chamber thickness for manual hemocytometers. Each sample chamber has two loading or vent ports 54, 55, one at each of the two opposing longitudinal ends of the elongated chamber. Overflow areas 56, 57, 58, 59 that are open at the top of the slide are positioned at each of the four corners of each sample chamber to accommodate excess sample and thereby insure that the sample chamber is properly filled with sample.

[0023] Since each sample chamber is closer to the upper plate 42 than to the lower plate 43, the slide functions best when properly inserted into the cell counter with the upper plate 42, and hence the thinnest optical window,

at the top. To ensure that the slide is inserted in this orientation, the slide is formed with notches 61, 62 in two diagonally opposing corners of the slide. The internal surfaces of the slot in the cell counter into which the slide is inserted to initiate the functions of the cell counter contains contour features that are complementary to these notches. The notches and complementary contours in the slot thereby prevent the user from inserting the slide upside down, *i. e.*, with the upper plate 42 at the bottom rather than the top. The symmetrical arrangement of the notches also complements the symmetrical arrangement of the two sample chambers and permits the slide to be inserted with either end first, while preventing the slide from being inserted in an inverted position (upside down). Since the slide is preferably a consumable item, it can thus be used for cell counting measurements on two independent samples at different times, and once both chambers have been used the slide can be disposed of and not used again.

[0024] Variations on the construction of the sample slide that still ensure proper orientation will be readily apparent to those skilled in the art. The arrangement, number, and shapes of the notches can thus be varied, as can the number of sample chambers and their locations relative to each other in the slide. The material of construction can vary widely and can be any material that can form an optically clear window, that is inert to the sample, and that is sufficiently rigid to be inserted into the cell counter. Poly(methyl methacrylate) and polycarbonate are examples of materials that can be used. Others will be readily apparent to those skilled in the art. Likewise, the Bonding of the plates can be accomplished by conventional means. Laser welding and ultrasonic welding are examples.

[0025] In the claims appended hereto, the terms "a" and "an" are intended to mean "one or more." The term "comprise" and variations thereof such as "comprises" and "comprising," when preceding the recitation of a step or an element, are intended to mean that the addition of further steps or elements is optional and not excluded.

Claims

1. An automated cell counter (11) comprising optical components, a sample mount (31) for receiving a sample slide (17), and a housing (12) retaining said optical components and sample mount (31), said optical components comprising a light source (22), a digital imaging sensor (25), and lenses (23, 26, 27, 29) positioned to direct light from said light source (22) to said digital imaging sensor (25) through said sample mount (31), said lenses (23, 26, 27, 29) comprising a negative lens (29) adjacent to said digital imaging sensor (25), and two achromat lenses (26, 27) of an achromat lens pair (24) separated by an aperture (28), the ratio of the focal lengths of said two achromat lenses (26, 27) being within a range of 1.5 to 6, said optical components arranged in a linear optical path and arranged such that when a sample slide (17) containing a cell suspension is mounted in said sample mount (31), an image of said cell suspension is projected onto said digital imaging sensor (25), said digital imaging sensor (25) comprising at least about 4,000,000 pixels and imaging a field of view at said sample mount (31) of at least 3 mm², each pixel being 2×2 μm or less in area, wherein said optical path is vertically oriented and said housing is mounted to a support base that is at least as wide as said housing (12), said housing (12) and support base (13) each occupying lateral areas of not more than 300 cm² and said optical path having a total height of not more than 20 cm.
2. The automated cell counter (11) of claim 1 wherein said lenses (23, 26, 27, 29) comprise a collimating lens (23) between said light source (22) and said sample slide (17).
3. The automated cell counter (11) of claim 1 wherein said digital sensor (25) contains from about 4,000,000 to about 10,000,000 pixels, each pixel is from about 0.5×0.5 μm to about 2×2 μm in area, and said imaging field is from about 3 mm² to about 10 mm².
4. The automated cell counter (11) of claim 1 wherein said digital imaging sensor (25) comprises means for generating contrast values between adjacent regions of said image, said sample mount (31) is of adjustable height along to said optical path, and said cell counter (11) further comprises means for automated adjustment of said height (32) in response to said contrast values until said image is focused.
5. The automated cell counter (11) of claim 4 further comprising a digital imager that counts cells in said field of view, and said digital imager and said automated adjustment of said height (32) of said sample mount (31) are initiated automatically upon insertion of a sample slide (17) in said sample mount (31).
6. The automated cell counter (11) of claim 1 wherein said digital imaging sensor (25) is flat, and said achromat lens pair (24) has a focus field at said digital imaging sensor (25).
7. The automated cell counter (11) of claim 1 wherein, when said support base (13) rests on a surface, said housing (12) is 30 cm or less in height and said sample mount (31) is at least 6 cm above said surface.
8. A method for counting cells in a cell suspension, said method comprising:
 - (a) placing an aliquot of said cell suspension in

an internal chamber of a sample slide (17) to form a film of said suspension in said chamber, (b) creating a digital image of a field of view of at least 3 mm² within said film on the automated cell counter (11) of claim 1 and (c) digitally counting said cells in said digital image.

9. The method of claim 8 wherein said digital imaging sensor (25) contains from about 4,000,000 to about 10,000,000 pixels, each pixel is from about 0.5×0.5 μm to about 2×2 μm in area, and said imaging field is from about 3 mm² to about 10 mm².
10. The method of claim 8 wherein step (b) is performed by a plurality of functions comprising automatic focusing of said digital image by moving said sample mount (31), detecting whether a vital stain is present in said aliquot, creating digital images of a plurality of focal planes within said internal chamber of said sample slide (17) when a vital stain is detected, and processing images formed on said digital imaging sensor (25) to determine a cell count in said aliquot, are all performed internally within said housing (12).
11. The method of claim 8 wherein step (c) comprises focusing said optical components on a plurality of planes within said internal chamber.
12. The method of claim 8, further comprising automatically generating contrast values between adjacent regions of said image, and automatically adjusting said sample mount (31) to a height along said optical path in response to said contrast values to focus said image, prior to step (b).
13. The method of claim 8, further comprising using a light source (22) that is spectrally limited to produce an image in said imaging sensor (25) that differentiates between live cells and dead cells.

Patentansprüche

1. Automatischer Zellzähler (11), der optische Komponenten, einen Probenhalter (31) zur Aufnahme eines Probenobjektträgers (17) und ein Gehäuse (12), in dem die optischen Komponenten und der Probenhalter (31) befestigt sind, umfasst, wobei die optischen Komponenten eine Lichtquelle (22), einen digitalen Bildgebungssensor (25) und Linsen (23, 26, 27, 29), die so positioniert sind, dass sie Licht aus der Lichtquelle (22) durch den Probenhalter (31) hindurch zu dem digitalen Bildgebungssensor (25) lenken, umfassen, wobei die Linsen (23, 26, 27, 29) eine Zerstreuungslinse (29), die dem digitalen Bildgebungssensor (25) am nächsten liegt, und zwei Achromatlinsen (26, 27) eines achromatischen Lin-

senpaares (24), die durch eine Blende (28) getrennt sind, umfassen, wobei das Verhältnis der Brennweiten der zwei Achromatlinsen (26, 27) innerhalb eines Bereichs von 1,5 bis 6 liegt, wobei die optischen Komponenten in einem linearen Lichtweg angeordnet sind und so angeordnet sind, dass dann, wenn ein Probenobjektträger (17), der eine Zellsuspension enthält, in dem Probenhalter (31) montiert ist, ein Bild der Zellsuspension auf den digitalen Bildgebungssensor (25) projiziert wird, wobei der digitale Bildgebungssensor (25) wenigstens etwa 4 000 000 Pixel umfasst und ein Gesichtsfeld an dem Probenhalter (31) von wenigstens 3 mm² abbildet, wobei jedes Pixel eine Fläche von 2 x 2 μm oder weniger hat, wobei der Lichtweg vertikal ausgerichtet ist und das Gehäuse auf einer Trägerbasis montiert ist, die wenigstens so breit wie das Gehäuse (12) ist, wobei das Gehäuse (12) und die Trägerbasis (13) jeweils seitliche Flächen von nicht mehr als 300 cm² einnehmen und der Lichtweg eine Gesamthöhe von nicht mehr als 20 cm aufweist.

2. Automatischer Zellzähler (11) gemäß Anspruch 1, wobei die Linsen (23, 26, 27, 29) einen Kollimator (23) zwischen der Lichtquelle (22) und dem Probenobjektträger (17) umfassen.
3. Automatischer Zellzähler (11) gemäß Anspruch 1, wobei der digitale Sensor (25) etwa 4 000 000 bis etwa 10 000 000 Pixel enthält, jedes Pixel eine Fläche von etwa 0,5 x 0,5 μm bis etwa 2 x 2 μm hat und das Gesichtsfeld etwa 3 mm² bis etwa 10 mm² beträgt.
4. Automatischer Zellzähler (11) gemäß Anspruch 1, wobei der digitale Bildgebungssensor (25) Einrichtungen zur Erzeugung von Kontrastwerten zwischen benachbarten Bereichen des Bildes umfasst, wobei der Probenhalter (31) eine entlang des Lichtwegs einstellbare Höhe aufweist und der Zellzähler (11) weiterhin Einrichtungen zur automatischen Einstellung der Höhe (32) als Reaktion auf die Kontrastwerte, bis das Bild fokussiert ist, umfasst.

5. Automatischer Zellzähler (11) gemäß Anspruch 4, weiterhin umfassend einen digitalen Bebilderer, der Zellen in dem Gesichtsfeld zählt, wobei der digitale Bebilderer und die automatische Einstellung der Höhe (32) des Probenhalters (31) beim Einsetzen eines Probenobjektträgers (17) in den Probenhalter (31) automatisch initiiert werden.
6. Automatischer Zellzähler (11) gemäß Anspruch 1, wobei der digitale Bildgebungssensor (25) flach ist und das achromatische Linsenpaar (24) einen Brennbereich auf dem digitalen Bildgebungssensor (25) aufweist.

7. Automatischer Zellzähler (11) gemäß Anspruch 1, wobei dann, wenn die Trägerbasis (13) auf einer Fläche aufliegt, das Gehäuse (12) eine Höhe von 30 cm oder weniger aufweist und sich der Probenhalter (31) wenigstens 6 cm oberhalb der Fläche befindet. 5
8. Verfahren zum Zählen von Zellen in einer Zellsuspension, wobei das Verfahren umfasst:
 - (a) Platzieren eines Aliquots der Zellsuspension in einer internen Kammer eines Probenobjektträgers (17) unter Bildung eines Films der Suspension in der Kammer; 10
 - (b) Schaffen eines digitalen Bildes eines Gesichtsfelds von wenigstens 3 mm² innerhalb des Films auf dem automatischen Zellzähler (11) gemäß Anspruch 1; und 15
 - (c) digitales Zählen der Zellen in dem digitalen Bild. 20
9. Verfahren gemäß Anspruch 8, wobei der digitale Bildgebungssensor (25) etwa 4 000 000 bis etwa 10 000 000 Pixel enthält, jedes Pixel eine Fläche von etwa 0,5 x 0,5 µm bis etwa 2 x 2 µm hat und das Gesichtsfeld etwa 3 mm² bis etwa 10 mm² beträgt. 25
10. Verfahren gemäß Anspruch 8, wobei Schritt (b) durch eine Vielzahl von Funktionen durchgeführt wird, umfassend das automatische Fokussieren des digitalen Bildes durch Bewegen des Probenhalters (31), das Nachweisen, ob in dem Aliquot ein Vitalfarbstoff vorhanden ist, das Schaffen digitaler Bilder einer Vielzahl von Brennebenen innerhalb der internen Kammer des Probenobjektträgers (17), wenn ein Vitalfarbstoff nachgewiesen wird, und das Verarbeiten der mit dem digitalen Bildgebungssensor (25) gebildeten Bilder, um eine Zellzahl in dem Aliquot zu bestimmen, die alle intern innerhalb des Gehäuses (12) durchgeführt werden. 30
11. Verfahren gemäß Anspruch 8, wobei Schritt (c) das Fokussieren der optischen Komponenten auf eine Vielzahl von Ebenen innerhalb der internen Kammer umfasst. 35
12. Verfahren gemäß Anspruch 8, weiterhin umfassend das automatische Erzeugen von Kontrastwerten zwischen benachbarten Bereichen des Bildes und das automatische Einstellen des Probenhalters (31) auf eine Höhe entlang des Lichtwegs als Reaktion auf die Kontrastwerte, um das Bild zu fokussieren, vor Schritt (b). 40
13. Verfahren gemäß Anspruch 8, weiterhin umfassend das Verwenden einer Lichtquelle (22), die spektral eingeschränkt ist, so dass in dem Bildgebungssensor (25) ein Bild entsteht, das zwischen lebenden Zellen und toten Zellen differenziert. 45

Revendications

1. Compteur automatique de cellules (11) comprenant des composants optiques, un support d'échantillon (31) destiné à recevoir une lame porte-échantillon (17), et un boîtier (12) retenant lesdits composants optiques et ledit support d'échantillon (31), lesdits composants optiques comprenant une source de lumière (22), un capteur d'image numérique (25), et des lentilles (23, 26, 27, 29) positionnées de façon à diriger la lumière de ladite source de lumière (22) vers ledit capteur d'image numérique (25) en traversant ledit support d'échantillon (31), lesdites lentilles (23, 26, 27, 29) comprenant une lentille négative (29) adjacente audit capteur d'image numérique (25), et deux lentilles achromatiques (26, 27) d'une paire de lentilles achromatiques (24) séparées par une ouverture (28), le rapport des distances focales desdites deux lentilles achromatiques (26, 27) étant compris dans une plage de 1,5 à 6, lesdits composants optiques étant disposés sur un chemin optique linéaire et étant disposés de façon que, quand une lame porte-échantillon (17) contenant une suspension de cellules est montée dans ledit support d'échantillon (31), une image de ladite suspension de cellules est projetée sur ledit capteur d'image numérique (25), ledit capteur d'image numérique (25) comprenant au moins environ 4 000 000 pixels et créant une image d'un champ de vision sur ledit support d'échantillon (31) d'au moins 3 mm², chaque pixel ayant une aire de 2 x 2 µm ou moins, ledit chemin optique étant orienté verticalement, et ledit boîtier étant monté sur une base support qui est au moins aussi large que ledit boîtier (12), ledit boîtier (12) et ladite base support (13) occupant chacun des zones latérales de pas plus de 300 cm², et ledit chemin optique ayant une hauteur totale de pas plus de 20 cm. 50
2. Compteur automatique de cellules (11) selon la revendication 1, dans lequel lesdites lentilles (23, 26, 27, 29) comprennent une lentille collimatrice (23) entre ladite source de lumière (22) et ladite lame porte-échantillon (17). 45
3. Compteur automatique de cellules (11) selon la revendication 1, dans lequel ledit capteur numérique (25) contient d'environ 4 000 000 à environ 10 000 000 pixels, chaque pixel ayant une aire d'environ 0,5 x 0,5 µm à environ 2 x 2 µm, et ledit champ d'imagerie étant d'environ 3 mm² à environ 10 mm². 50
4. Compteur automatique de cellules (11) selon la revendication 1, dans lequel ledit capteur d'image numérique (25) comprend un moyen de génération de valeurs de contraste entre des régions adjacentes de ladite image, ledit support d'échantillon (31) ayant une hauteur réglable le long dudit chemin optique, 55

et ledit compteur de cellules (11) comprenant en outre un moyen pour le réglage automatique de ladite hauteur (32) en réponse auxdites valeurs de contraste jusqu'à mise au point de ladite image.

5. Compteur automatique de cellules (11) selon la revendication 4, comprenant en outre un imageur numérique qui compte les cellules dans ledit champ de vision, et ledit imageur numérique et ledit réglage automatique de ladite hauteur (32) dudit support d'échantillon (31) étant déclenchés automatiquement lors de l'insertion d'une lame porte-échantillon (17) dans ledit support d'échantillon (31). 5 10
6. Compteur automatique de cellules (11) selon la revendication 1, dans lequel ledit capteur d'image numérique (25) est plat, et ladite paire de lentilles achromatiques (24) présente un champ focal au niveau dudit capteur d'image numérique (25). 15 20
7. Compteur automatique de cellules (11) selon la revendication 1, dans lequel, quand ladite base support (13) repose sur une surface, ledit boîtier (12) présente une hauteur de 30 cm ou moins, et ledit support d'échantillon (31) se trouve à au moins 6 cm au-dessus de ladite surface. 25
8. Procédé de comptage de cellules dans une suspension de cellules, ledit procédé comprenant : 30
 - (a) le placement d'une aliquote de ladite suspension de cellules dans une chambre interne d'une lame porte-échantillon (17) pour former un film de ladite suspension dans ladite chambre, 35
 - (b) la création d'une image numérique d'un champ de vision d'au moins 3 mm² à l'intérieur dudit film sur le compteur automatique de cellules (11) selon la revendication 1, et
 - c) le comptage numérique desdites cellules dans ladite image numérique. 40
9. Procédé selon la revendication 8, dans lequel ledit capteur d'image numérique (25) contient d'environ 4 000 000 à environ 10 000 000 pixels, chaque pixel a une aire d'environ 0,5 x 0,5 µm à environ 2 x 2 µm, et ledit champ d'imagerie est d'environ 3 mm² à environ 10 mm². 45
10. Procédé selon la revendication 8, dans lequel l'étape (b) est mise en oeuvre par une pluralité de fonctions comprenant la mise au point automatique de ladite image numérique par déplacement dudit support d'échantillon (31), la détection de si une coloration vitale est présente dans ladite aliquote, la création d'images numériques d'une pluralité de plans focaux à l'intérieur de ladite chambre interne de ladite lame porte-échantillon (17) quand une coloration vitale est 50 55

détectée, et le traitement d'images formées sur ledit capteur d'image numérique (25) pour déterminer un comptage cellulaire dans ladite aliquote, toutes étant mises en oeuvre à l'intérieur dudit boîtier (12).

11. Procédé selon la revendication 8, dans lequel l'étape (c) comprend la mise au point desdits composants optiques sur une pluralité de plans à l'intérieur de ladite chambre interne.
12. Procédé selon la revendication 8, comprenant en outre la génération automatique de valeurs de contraste entre des régions adjacentes de ladite image, et le réglage automatique dudit support d'échantillon (31) à une hauteur le long dudit chemin optique en réponse auxdites valeurs de contraste pour mettre au point ladite image, avant l'étape (b).
13. Procédé selon la revendication 8, comprenant en outre l'utilisation d'une source de lumière (22) qui est spectralement limitée pour produire une image dans ledit capteur d'image (25), qui fait la distinction entre des cellules vivantes et des cellules mortes.

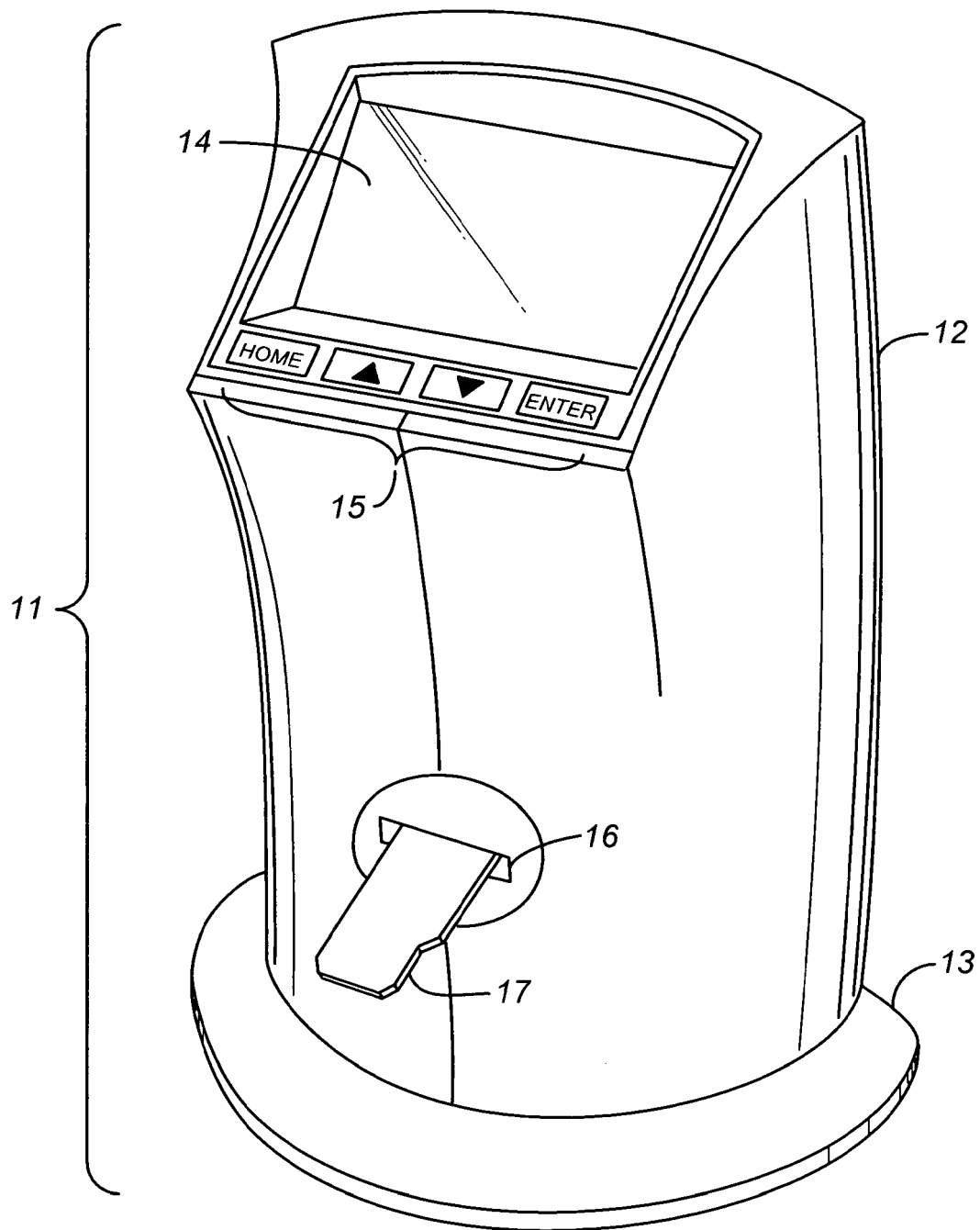


FIG. 1

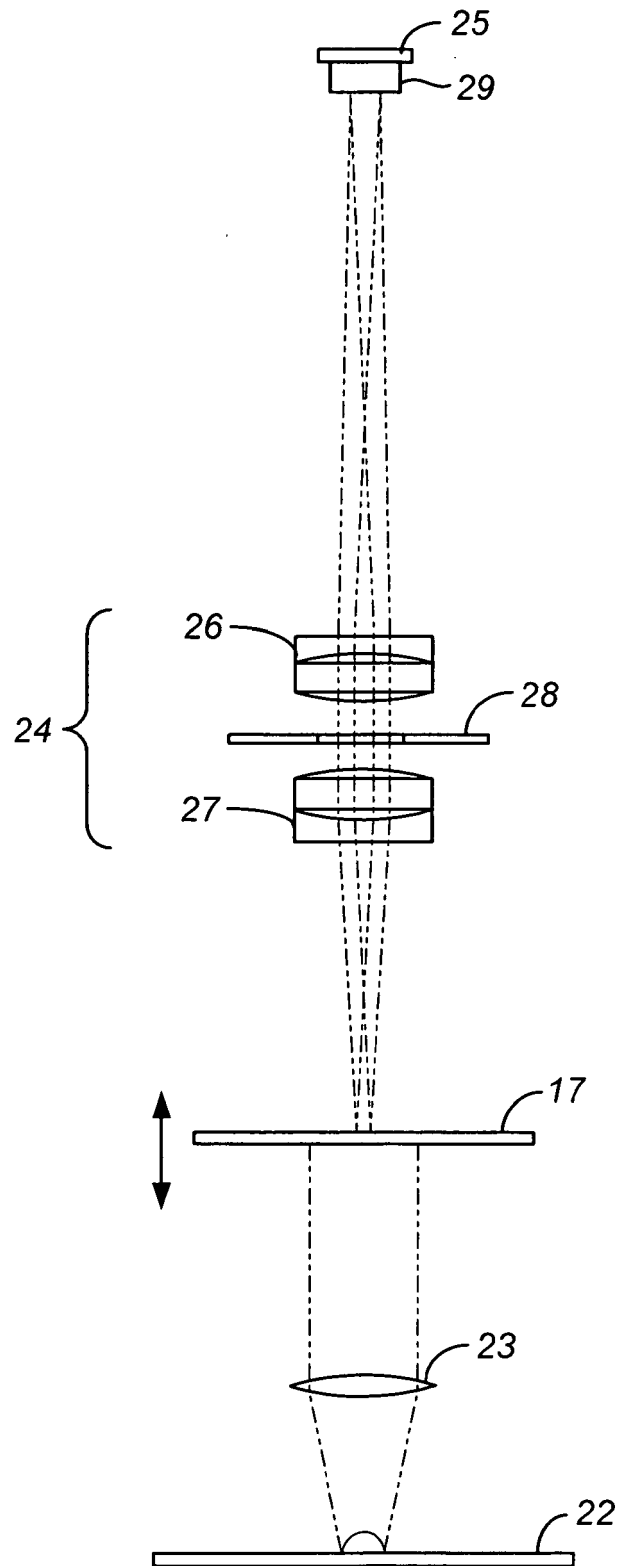


FIG. 2

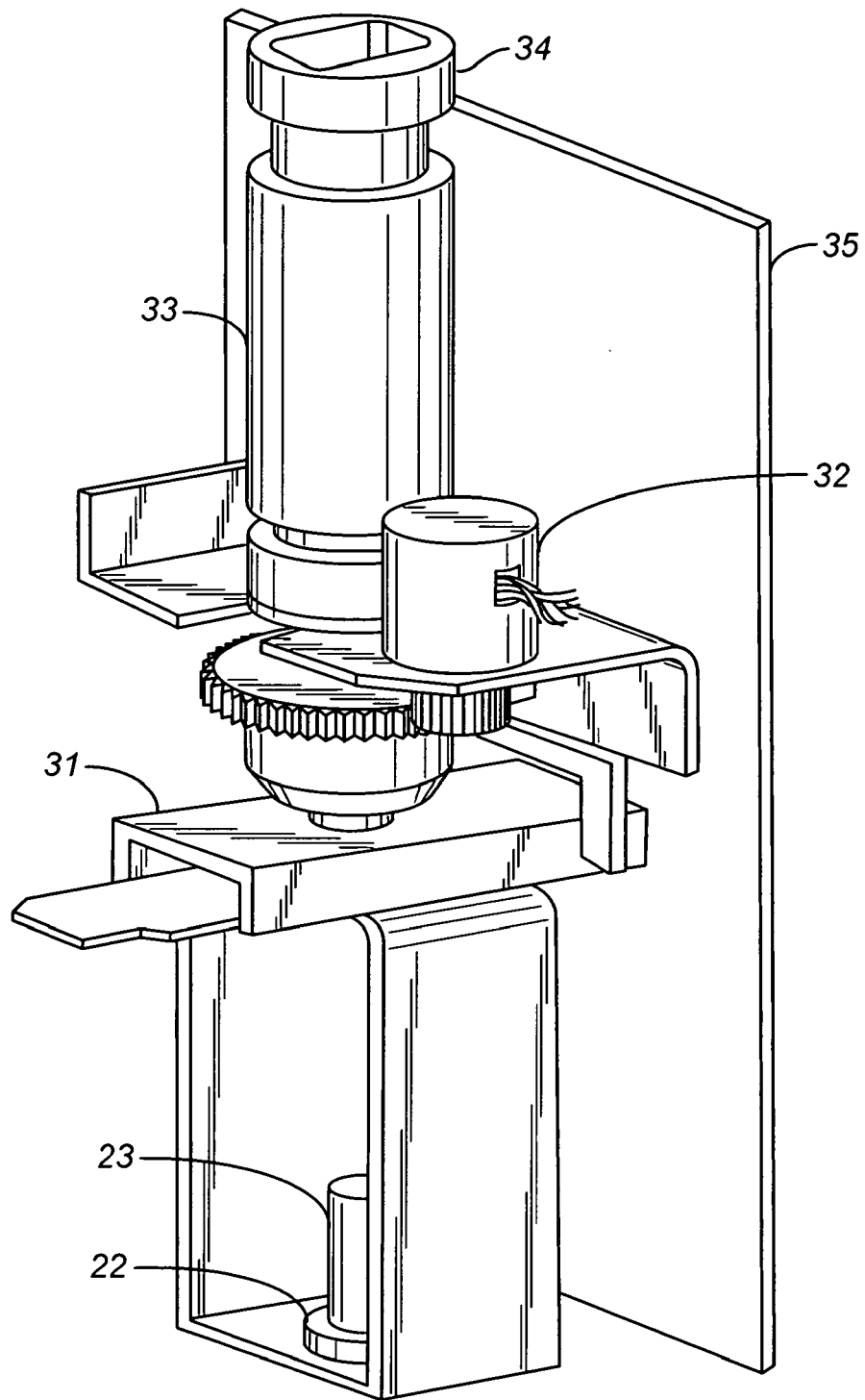


FIG. 3

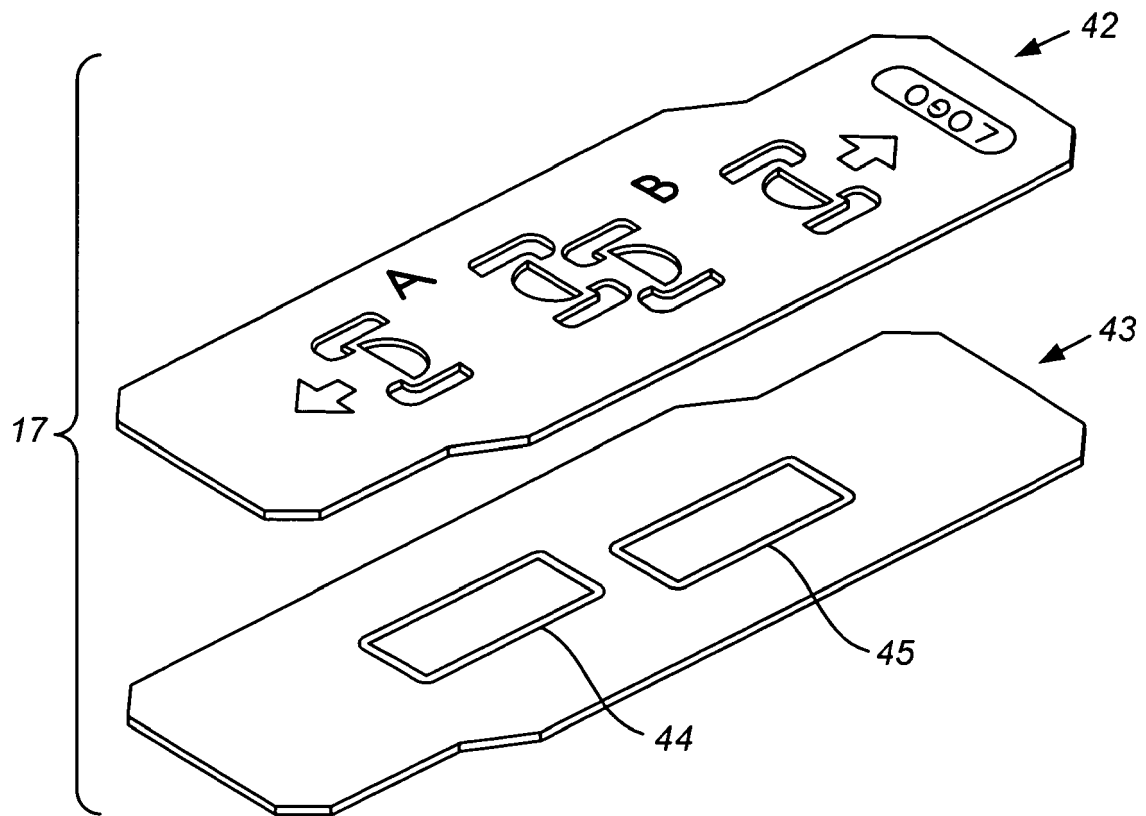
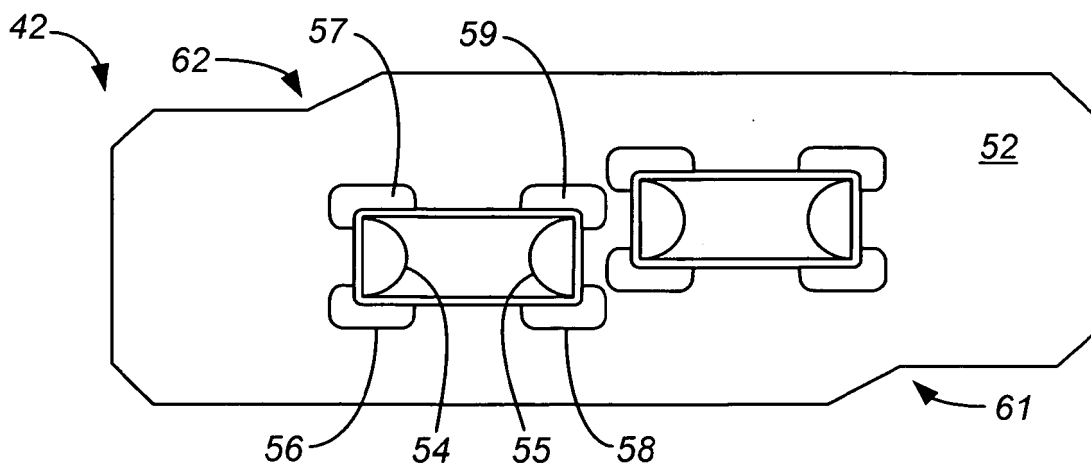
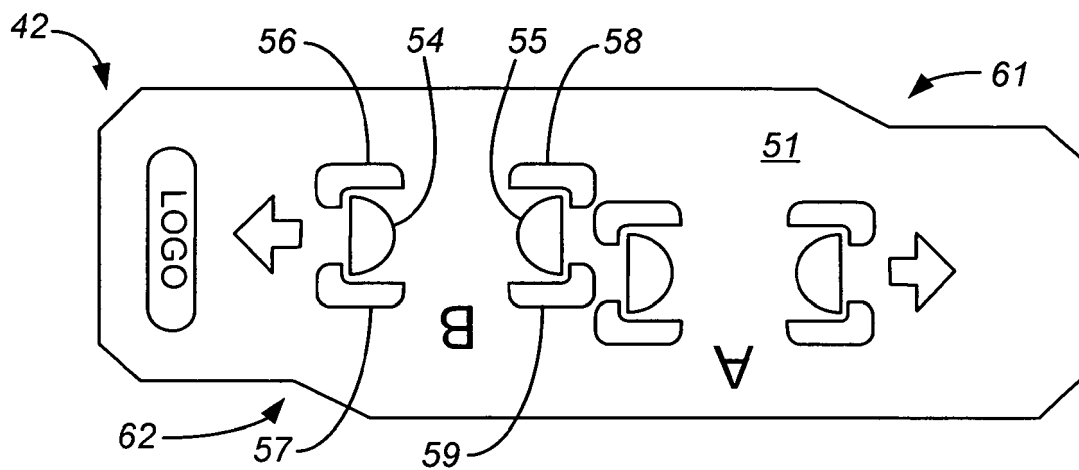


FIG. 4



REFERENCES CITED IN THE DESCRIPTION

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